LISTING OF CLAIMS

Claims 1-10 (CANCELED)

- 5 11- (NEW) A solid orodispersible pharmaceutical composition comprising:
 - granules consisting of co-dried lactose and starch, and
 - perindopril or a pharmaceutically acceptable salt thereof.
- 12- (NEW) A composition according to claim 11, wherein the composition disintegrates in the mouth in less than three minutes.
 - 13- (NEW) A composition according to claim 12, wherein the composition disintegrates in the mouth in less than one minute.
- 14- (NEW) A composition according to claim 11, comprising, in relation to the total weight of the composition:
 - from 85 % to 99 % by weight of granules consisting of co-dried lactose and starch, and
 - from 0.1 % to 10 % by weight of perindopril or a pharmaceutically acceptable salt thereof.
- 15- (NEW) A composition according to claim 14, comprising from 0.5 % to 6 % by weight of perindopril or a pharmaceutically acceptable salt thereof.
 - 16- (NEW) A composition according to claim 11, further comprising one or more lubricants, a flow agent and, optionally, a sweetening agent.
- 17- (NEW) A composition according to claim 11, wherein the composition is in the form of a tablet.
 - 18- (NEW) A tablet according to claim 17, wherein the tablet is obtained by direct compression.

- 19- (NEW) A tablet according to claim 18, wherein the tablet has a hardness from 5 to 50 Newtons.
- 20- (NEW) A tablet according to claim 19, wherein the tablet has a hardness from 10 to 20 Newtons.
- 21- (NEW) A process for the manufacture of solid orodispersible compositions of perindopril, or a pharmaceutically acceptable salt thereof, which disintegrate in the mouth in less than three minutes, wherein the perindopril or a pharmaceutically acceptable salt thereof is mixed with granules consisting of co-dried lactose and starch.
- 22- (NEW) A process for the manufacture of solid orodispersible compositions of perindopril, or a pharmaceutically acceptable salt thereof, which disintegrate in the mouth in less than one minute, wherein the perindopril or a pharmaceutically acceptable salt thereof is mixed with granules consisting of co-dried lactose and starch.
- 23- (NEW) A method for treating a living animal body including a human afflicted with a condition selected from arterial hypertension and heart failure comprising the step of administering to the living animal body including a human a composition according to claim 11 which is effective for alleviation of the condition.